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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
09/451,641	11/30/1999	Danchen Gao	C-3169/1/US	9327	
26648	7590 05/17/2005		EXAMINER		
PHARMACIA CORPORATION			TRAN, SUSAN T		
GLOBAL PA	TENT DEPARTMENT				
POST OFFICE BOX 1027			ART UNIT	PAPER NUMBER	
ST. LOUIS, N	MO 63006		1615		
			DATE MAIL ED. 05/17/000	_	

Please find below and/or attached an Office communication concerning this application or proceeding.

PTOL-326 (Rev. 1-04)	Office Action Summa	гу	Part of Paper No./Mail Date 050405	. /
Attachment(s) 1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-3) Information Disclosure Statement(s) (PTO-1449 or Paper No(s)/Mail Date 02/14/05. 2/24/05	•	4) Interview Summary Paper No(s)/Mail Da 5) Notice of Informal P 6) Other:		
12) Acknowledgment is made of a claim f a) All b) Some * c) None of: 1. Certified copies of the priority of 2. Certified copies of the priority of 3. Copies of the certified copies of application from the Internation * See the attached detailed Office action	documents have bee documents have bee of the priority documental Bureau (PCT Rule	n received. n received in Applicati ents have been receive e 17.2(a)).	on No ed in this National Stage	
Priority under 35 U.S.C. § 119				
9) The specification is objected to by the 10) The drawing(s) filed on is/are: Applicant may not request that any object Replacement drawing sheet(s) including 11) The oath or declaration is objected to	a) accepted or b) tion to the drawing(s) be the correction is require	e held in abeyance. See ed if the drawing(s) is obj	e 37 CFR 1.85(a). jected to. See 37 CFR 1.121(d).	
8) Claim(s) are subject to restrict	tion and/or election re	equirement.		
5) ☐ Claim(s) is/are allowed. 6) ☑ Claim(s) <u>1, 2, 4-10,12-50,72-75,84 allowed.</u> 7) ☑ Claim(s) <u>3</u> is/are objected to.	<u>nd 86-90</u> is/are rejec	ed.		
4) Claim(s) <u>1-10,12-50,72-75,84 and 86</u> 4a) Of the above claim(s) is/ar				
Disposition of Claims	S 00 is/ara panding in	the application		
closed in accordance with the practic	e under <i>Ex parte Qu</i>	<i>ayle</i> , 1935 C.D. 11, 45	53 O.G. 213.	
3) Since this application is in condition f	•			
1) Responsive to communication(s) file2a) This action is FINAL.	b on <u>24 February 200</u> b)⊡ This action is n			
Status				
A SHORTENED STATUTORY PERIOD FO THE MAILING DATE OF THIS COMMUNION. - Extensions of time may be available under the provisions of after SIX (6) MONTHS from the mailing date of this commu- lif the period for reply specified above is less than thirty (30). - If NO period for reply is specified above, the maximum states and the second period for reply within the set or extended period for reply within the set or ext	CATION. of 37 CFR 1.136(a). In no evenuication. b) days, a reply within the statututory period will apply and wiwill, by statute, cause the apple.	ent, however, may a reply be tim story minimum of thirty (30) days ll expire SIX (6) MONTHS from ication to become ABANDONE	nely filed s will be considered timely. the mailing date of this communication. D (35 U.S.C. § 133).	
The MAILING DATE of this communic	cation appears on the	cover sheet with the c	orrespondence address	
	Susan T.	ran (1615	
Office Action Summary	09/451,64 Examiner		GAO ET AL. Art Unit	
	Application		Applicant(s)	

TH

DETAILED ACTION

Receipt is acknowledged of applicant's Petition to Withdraw from Issue, Request for Continued Examination, and Information Disclosure Statement filed 02/24/05.

Continued Examination Under 37 CFR 1.114

A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after allowance or after an Office action under *Ex Parte Quayle*, 25 USPQ 74, 453 O.G. 213 (Comm'r Pat. 1935). Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, prosecution in this application has been reopened pursuant to 37 CFR 1.114. Applicant's submission filed on 02/24/05 has been entered.

Information Disclosure Statement

The information disclosure statement (IDS) submitted on 02/24/05 was filed after the mailing date of the allowance on 11/21/03. The submission is in compliance with the provisions of 37 CFR 1.97. Accordingly, the information disclosure statement is being considered by the examiner.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

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(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

Claims 1, 2, 4-10, 12-50, 72-75, 84 and 86-90 are rejected under 35 U.S.C. 103(a) as being unpatentable over AAPS Annual Meeting Contributed Papers Abstracts (AAPS), in view of Black EP 0 863 134.

AAPS teaches a celecoxib (Cox-2 inhibitor) formulation that exhibits a C_{max} values of 1527 and 1077 ng/mL, and a T_{max} of 1.9 hours (see page D32).

AAPS does not teach the use of excipients in the formulation. However, the use of excipients in oral formulations is well known in pharmaceutical art. To be more specific, Black teaches a compound useful as a Cox-2 inhibitor for pain relief, fever and inflammation of a variety symptoms disclosed on page 3, lines 29-36. The compound can be administered orally in the form of tablets, troches, lozenges, or capsules (page 4, lines 1-12). The tablets comprising active ingredient in admixture with excipients, e.g., diluents, disintegrants, binding agents, wetting agents, and surfactant (page 4, lines 15-38). The active agent present in an amount of 10 to 250 mg, and carrier material may vary from about 5 to about 95% (page 5, lines 39-58). The dosage can be administered once or twice a day, and will provide effective T_{1/2} over a 24 hours period (page 5, lines 22-27). Example 2 discloses the amount of excipients use in a tablet. Thus, it would have been obvious for one of ordinary skill in the art to prepare the formulation of AAPS using excipient in view of the teachings of Black to obtain the claimed invention, because Black teaches oral dosage of Cox-2 inhibitor, and because

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AAPS teaches orally administering celecoxib in fine suspension and capsule forms having the claimed C_{max} and T_{max} values.

It is noted that AAPS does not expressly teach the particle size distribution, however, the burden is shifted to applicant to show that the formulation of AAPS does not have the claimed particle size distribution, as well as the detrimental effect and/or unexpected results over the particle size distribution, because AAPS teaches the oral formulation of celecoxib having the claimed C_{max} and T_{max} values.

Conclusion

All claims are drawn to the same invention claimed in the application prior to the entry of the submission under 37 CFR 1.114 and could have been finally rejected on the grounds and art of record in the next Office action if they had been entered in the application prior to entry under 37 CFR 1.114. Accordingly, THIS ACTION IS MADE **FINAL** even though it is a first action after the filing of a request for continued examination and the submission under 37 CFR 1.114. See MPEP § 706.07(b). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of

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the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

Claims Allowable

Claim 3 is objected to as being dependent upon a rejected base claim, but would be allowable if rewritten in independent form including all of the limitations of the base claim and any intervening claims.

Correspondence

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Susan T. Tran whose telephone number is (571) 272-0606. The examiner can normally be reached on M-R from 6:00 am to 4:30 pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Thurman K. Page, can be reached at (571) 272-0602. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should

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you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

THURMAN K. PAGE
SUPERVISORY PATENT EXAMINER
TECHNOLOGY CENTER 1600